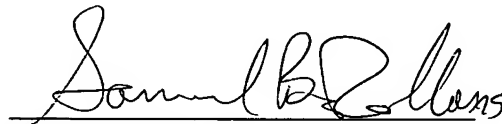


all claims and entry of new claims 72-75, which are identical to the claims 39-42 in the originally filed application.

The Examiner is invited to contact the undersigned attorney regarding any questions related to this application.

Respectfully submitted,

Date: 10/15/03


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AMINATION PROCESS

Field of the Invention

[0001] The present invention belongs to the field of synthetic organic chemistry. In particular, it relates to a process for forming carbon-nitrogen bonds, particularly vinyl-nitrogen bonds and aryl-nitrogen bonds.

Background of the Invention

[0002] The development of methods that effect the formation of carbon-nitrogen bonds is a challenge of broad interest due to the prevalence of the aniline subunit within many biologically-active natural products and medicinal agents (See, *e.g.*, The Alkaloids: Chemistry and Biology, Cordell, G.A., Ed.; Academic: San Diego, 1998, Vol. 50). Significantly, indole alkaloid syntheses frequently commence from one of many available aniline or indole derivatives. Although this strategy is inherently limited, its popularity is perhaps due to the fact that there are few methods available for forming aryl-nitrogen bonds. Moreover, existing technology generally fails to offer the mild conditions necessary for the highest degree of chemoselectivity. Transition-metal mediated aryl amination generally requires the use of a basic additive to promote the coupling process. Moreover, transition metal ligand selection is based upon consideration of the individual electronic nature of the aromatic halide (or triflate, etc.) and amine components.

[0003] Similar, but fewer, methods are available for vinyl amination. Again, the need for basic and/or nucleophilic addends limits the substrate generally of these methodologies. Metal-mediated alkyne amination offers alternative access to the products of vinyl amination (enamines), but functional group tolerance is attenuated further still. Notwithstanding the technological limitations, the products of these transformations are valuable both as synthetic intermediates and as targets themselves. Of particular importance is the pyrrolidine heterocyclic class and its oxidized variants (dihydropyrrole and pyrrole). For example, 2-carboxy pyrrolidine is also known as proline, an α -amino acid prevalent in biopeptides. The pyrrolidine backbone is also found in medicinal agents (*i.e.*, drugs) and numerous classes of natural products displaying a range of biological activity.

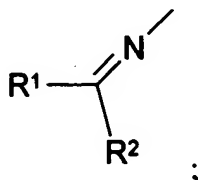
Summary of the Invention

[0004] The present invention provides methodology for carbon-nitrogen bond formation via vinyl or aryl amination. In the process of the invention, an sp^2 hybridized radical is reacted with an azomethine moiety to form dihydropyrrole, 2-methylenopyrrolidine, and indoline compounds. The methodology provides a facile process for the synthesis of compounds having the pyrrolidine or indoline subunit and is especially advantageous for compounds having acid or base labile functional groups and/or is comprised of chiral centers susceptible to acid/base epimerization.

Detailed Description of the Invention

[0005] In a first aspect, the present invention provides a process for forming an intramolecular carbon-nitrogen bond which comprises reacting an sp^2 hybridized carbon radical moiety with an azomethine moiety in the presence of a hydrogen atom donor, wherein said azomethine moiety possesses at least one radical stabilizing group, and the azomethine carbon is in the ketone oxidation state or higher. In this regard, suitable radical stabilizing groups are known in the art and include but are not limited to the following: phenyl, vinyl, trifluormethyl, carbonyl, and the like. In a preferred embodiment, the azomethine carbon will be in the ketone oxidation state and will be bonded by groups such as hydrocarbyl, substituted hydrocarbyl, aryl, and heteroaryl, provided that the atom bonded between such groups and the azomethine carbon is a carbon atom.

[0006] As used herein, the term "azomethine" preferably refers to the subunit having the Formula



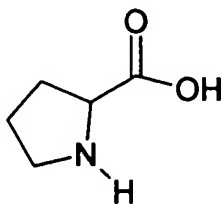
wherein R¹ and R² are as defined herein.

[0007] In this process, the sp^2 hybridized radical can be formed using any number of methodologies, including but not limited to preparation of the desired carbon radical moiety by photolysis, thermal cleavage, or by homolytic transmetalation in the presence of a free radical

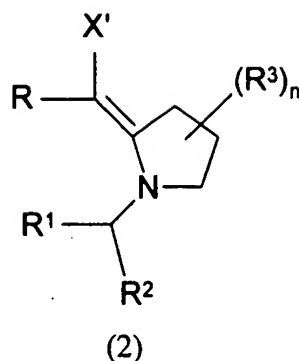
initiator compound. In the latter case, the substituted carbon moiety will thus preferably be substituted by a suitable radical leaving group such as a halogen atom. The term "free radical initiator" compound is any compound which is capable of facilitation of a free radical reaction via a homolytic mechanism. Examples include azonitrile compounds such as 2,2'-azobisisobutyronitrile (AIBN); peroxides; and the like. The carbon radical may also be produced via a prior homolytic reaction, including but not limited to a radical addition to an olefin or the thermal cyclization of an ene-diyne moiety.

[0008] Preferred "hydrogen atom donor" compounds include reactants or species which can be generated *in situ* which provide a hydrogen atom. Examples of suitable hydrogen donor compounds include organostannane hydrides, organosilyl silanes, organogermanium hydrides, 1,4-cyclohexadiene, γ -terpinene, thiols, selenol, and the like. Examples of organostannane hydrides include compounds of the Formula $(X')_3\text{Sn-H}$, wherein X' is an alkyl group, preferably a $\text{C}_1\text{-C}_6$ alkyl group, aryl group, or a fluoros derivative thereof. Alternatively, such a compound can be generated *in situ*; for example, hexamethylditin can be photolyzed to provide the same tin radical as tri-n-butyl tin hydride plus a free radical initiator compound. Examples of alkylsilylsilanes include tris(trimethylsilyl)silane, triethylsilane, and the like.

[0009] The process of the invention is particularly well suited for the preparation of various pyrrolidine compounds. One such compound is proline (or its derivatives):



[0010] Thus, in a second aspect, the present invention provides a process for preparing a compound of Formula (2)

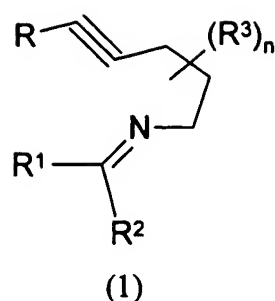


wherein each R is independently selected from the group consisting of hydrogen, hydrocarbyl, substituted hydrocarbyl, aryl, heteroaryl, substituted aryl, substituted heteroaryl, heteroatom connected hydrocarbyl, heteroatom connected substituted hydrocarbyl, heteroatom connected aryl, heteroatom connected heteroaryl, heteroatom connected substituted aryl, heteroatom connected substituted heteroaryl, a group of the formula $-C(O)R^1$, a group of the formula $-O-R^1$, a group of the formula $-NHR^1$, a group of the formula $-N(R^1)_2$, a group of the formula $-Sn(R^1)_3$, and a group of the formula $-Si(R^1)_3$;

wherein the R^1 and R^2 groups are independently selected from the group consisting of aryl, heteroaryl, hydrocarbyl, substituted aryl, substituted heteroaryl, and substituted hydrocarbyl; provided that said groups are bonded via a carbon atom;

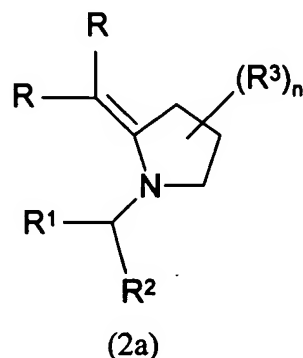
each R^3 is independently selected from aryl; heteroaryl; hydrocarbyl; substituted aryl; substituted heteroaryl; substituted hydrocarbyl; heteroatom connected aryl; heteroatom connected hydrocarbyl; heteroatom connected substituted hydrocarbyl; heteroatom connected heteroaryl; heteroatom connected substituted aryl; halo, preferably fluoro or chloro, most preferably fluoro; amino; cyano; hydroxy; carboxy; a group of the formula $-C(O)O-C_1-C_8$ alkyl; a group of the formula $-C(O)R^1$; a group of the formula $-O-R^1$; a group of the formula $-NHR^1$; a group of the formula $-N(R^1)_2$; C_1-C_8 alkoxy; C_1-C_8 alkylthio; and oxo (*i.e.*, an in-line carbonyl group wherein the oxygen is doubly bonded with the carbon atom to which R^3 is attached, in which case n will of course be 1); or two R^3 groups taken together can form a divalent hydrocarbyl, substituted hydrocarbyl, or be bonded directly to a heteroatom such as oxygen, nitrogen, or sulfur; and n is from 0 to 6;

which comprises contacting a compound of Formula (1)



with a free radical initiator in the presence of a hydrogen atom donor, wherein R, R¹, R², R³, and n are as defined above.

[0011] In a third aspect, there is provided a process for preparing compounds of the formula

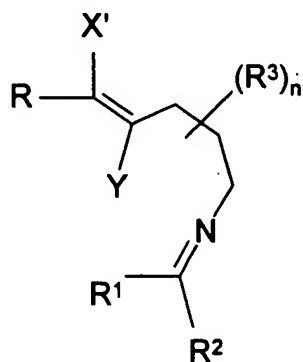


wherein each R is independently selected from the group consisting of hydrogen, hydrocarbyl, substituted hydrocarbyl, aryl, heteroaryl, substituted aryl, substituted heteroaryl, heteroatom connected hydrocarbyl, heteroatom connected substituted hydrocarbyl, heteroatom connected aryl, heteroatom connected heteroaryl, heteroatom connected substituted aryl, heteroatom connected substituted heteroaryl, a group of the formula $-C(O)R^1$, a group of the formula $-O-R^1$, a group of the formula $-NHR^1$, a group of the formula $-N(R^1)_2$, a group of the formula $-Sn(R^1)_3$, and a group of the formula $-Si(R^1)_3$;

wherein the R¹ and R² groups are independently selected from the group consisting of aryl, heteroaryl, hydrocarbyl, substituted aryl, substituted heteroaryl, and substituted hydrocarbyl; provided that said groups are bonded via a carbon atom;

each R^3 is independently selected from aryl; heteroaryl; hydrocarbyl; substituted aryl; substituted heteroaryl; substituted hydrocarbyl; heteroatom connected aryl; heteroatom connected hydrocarbyl; heteroatom connected substituted hydrocarbyl; heteroatom connected heteroaryl; heteroatom connected substituted aryl; amino; halo; cyano; hydroxy; carboxy; a group of the formula $-C(O)O-C_1-C_8$ alkyl; a group of the formula $-C(O)R^1$; a group of the formula $-O-R^1$; a group of the formula $-NHR^1$; a group of the formula $-N(R^1)_2$; C_1-C_8 alkoxy; C_1-C_8 alkylthio; and oxo; or two R^3 groups taken together can form a divalent hydrocarbyl, substituted hydrocarbyl, or be bonded directly to a heteroatom selected from oxygen, nitrogen, or sulfur; and each n is 0, 1 or 2;

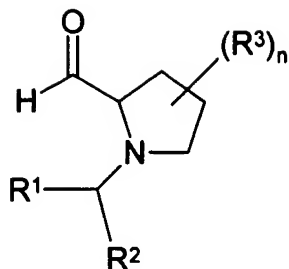
which comprises contacting a compound of the formula



with a free radical initiator in the presence of a hydrogen atom donor, wherein Y is a radical leaving group, and X' is a group selected from C_1-C_6 alkyl, aryl, or a fluorous derivative thereof.

[0012] The compounds of Formula (2) and (2a) can thus be derivatized to form proline by epoxidation and acid-catalyzed rearrangement to the amino aldehyde. Oxidation and deprotection of the amine provides proline; in this fashion, proline and other proline-subunit containing compounds may be synthesized using intermediates of Formula (2). Thus, as a fourth aspect of the invention, there is provided the second aspect of the invention as set forth above, further comprising the steps:

- (a) epoxidation, followed by acid catalyzed rearrangement to afford an amino aldehyde, of the formula

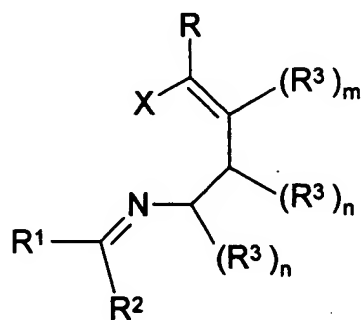


followed by

- (b) treatment of the resulting aldehyde with a suitable inorganic oxidizing agent,
- (c) followed by deprotection of the nitrogen to provide proline, wherein R^1 , R^2 , R^3 , and n are as defined above.

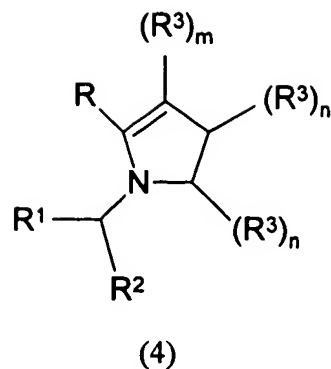
[0013] In this regard, suitable inorganic oxidizing agents include chromic acid, and suitable methodologies for removal of the R^1 and R^2 groups (*i.e.*, deprotection of the nitrogen) include conventional hydrogenation utilizing, for example, H_2 , Pd/C, HCO_2H or HCl, in diethyl ether followed by appropriate workup.

[0014] As an alternative to the alkyne starting material of Formula (1) above, one may utilize the corresponding vinyl halide of Formula (3)



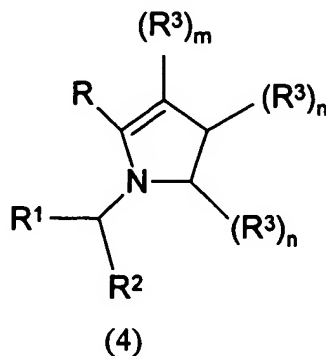
(3)

wherein R , R^1 , R^2 , and R^3 , and n are as defined above, m is zero or one (in which case, the carbon to which R^3 is attached will be substituted by hydrogen), and X is a halide such as bromo, iodo, or chloro, which provides access to dihydropyrrolidine ring systems having the formula



[0015] In the above formula (3), it will be understood that the $(R^3)_n$ group represents that there may be one, two or no such R^3 groups at the 4 and 5 positions of the dihydropyrrole ring. It will also be understood that either olefin stereoisomer may be utilized in the above method insofar as the vinyl radical stereoisomers epimerize rapidly and only one leads to the cyclized product.

[0016] Thus, in a fifth aspect, the present invention provides a process for preparing a compound of the Formula (4)

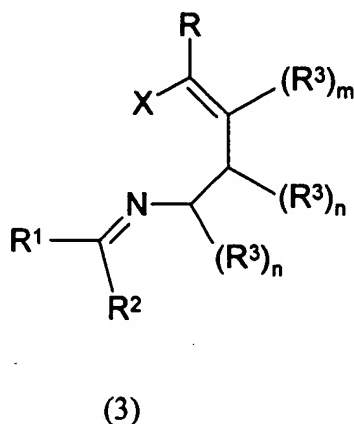


wherein each R is independently selected from the group consisting of hydrogen, hydrocarbyl, substituted hydrocarbyl, aryl, heteroaryl, substituted aryl, substituted heteroaryl, heteroatom connected hydrocarbyl, heteroatom connected substituted hydrocarbyl, heteroatom connected aryl, heteroatom connected heteroaryl, heteroatom connected substituted aryl, heteroatom connected substituted heteroaryl, a group of the formula $-C(O)R^1$, a group of the formula $-O-R^1$, a group of the formula $-NHR^1$, a group of the formula $-N(R^1)_2$, a group of the formula $-Sn(R^1)_3$, and a group of the formula $-Si(R^1)_3$;

wherein the R^1 and R^2 groups are independently selected from the group consisting of aryl, heteroaryl, hydrocarbyl, substituted aryl, substituted heteroaryl, and substituted hydrocarbyl; provided that said groups are bonded via a carbon atom;

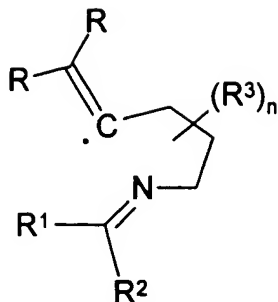
each R^3 is independently selected from aryl; heteroaryl; hydrocarbyl; substituted aryl; substituted heteroaryl; substituted hydrocarbyl; heteroatom connected aryl; heteroatom connected hydrocarbyl; heteroatom connected substituted hydrocarbyl; heteroatom connected heteroaryl; heteroatom connected substituted aryl; amino; halo; cyano; hydroxy; carboxy; a group of the formula $-C(O)O-C_1-C_8$ alkyl; a group of the formula $-C(O)R^1$; a group of the formula $-O-R^1$; a group of the formula $-NHR^1$; a group of the formula $-N(R^1)_2$; C_1-C_8 alkoxy; C_1-C_8 alkylthio; and oxo; or two R^3 groups taken together can form a divalent hydrocarbyl, substituted hydrocarbyl, or be bonded directly to a heteroatom selected from oxygen, nitrogen, or sulfur; m is 0 or 1; and each n is 0, 1 or 2;

which comprises contacting a compound of the Formula (3)



wherein R, R^1 , R^2 , and R^3 , and n are as defined for Formula (4), and X is a halide, with a free radical initiator in the presence of a hydrogen atom donor.

[0017] In the process of this aspect of the invention, the desired radical may be generated *in situ* using methodology described above with regard to the first aspect. Thus, in a further embodiment, there is provided the free radical intermediate of the Formula



wherein R, R¹, R², R³ and n are as defined above,
which is useful as an intermediate in the preparation of the compounds of Formula (2).

[0018] The advantages of the methodology of this invention include the readily-available amino-alkyne substrates of Formula (1) as depicted above, as well as the use of radical intermediates and reaction conditions which are pH neutral. Thus, no acid or base addends are necessary to effect the desired transformation. This feature is especially advantageous when the desired pyrrolidine compound possesses acid or base labile functional groups and/or is comprised of chiral center(s) susceptible to acid/base-catalyzed epimerization. Moreover, there are no electronic restrictions on the substrates, and thus acid- and base-sensitive substrates that are electron rich, electron neutral, or electron deficient may be utilized. This addition of vinyl radicals to the nitrogen of an azomethine moiety may result in excess of 10:1, preferably 20:1 regioselectivity for the desired carbon- nitrogen bond formation. In contrast to other methodologies for amination, the active nitrogen-containing component in each case is an azomethine, and the pi-bond is its reactive feature (as opposed to a sigma N-H bond).

[0019] In the above Formulae (1) and (2), it will be understood that the indeterminate placement of the R³ group(s) denotes that there may be none or up to 6 of such groups at the carbon atoms α , β , and γ to the nitrogen atom. In this regard, the possible substituents from which R³, as well as R¹ and R² may be selected is not particularly limited, so long as such groups do not contain species which have a deleterious effect on the desired reaction or otherwise serve to quench the sp²-hybridized carbon radical prematurely.

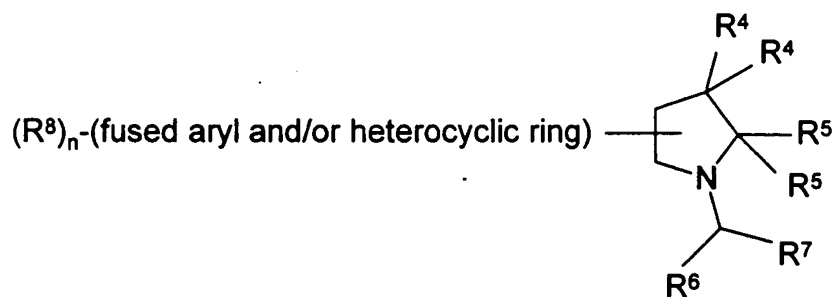
[0020] Further examples of starting materials (1) of the present invention include (Z) or (E)-vinyl halides. In addition, the process of the present invention may be conducted on substrates which are attached to a solid support via any carbon of the required structural features. In addition, the process of the present invention may be utilized in the construction of combinatorial

libraries of compounds. Utilizing the methodology of the present invention, enantiomerically enriched starting materials may be utilized to prepare enantiomerically enriched final products.

[0021] The methodology of the present invention is also useful in the context of aryl amination. Thus, in a third aspect, the present invention provides a process for forming a carbon-nitrogen bond, wherein said carbon is part of an aryl or heteroaryl ring, which comprises reacting an aryl or heteroaryl radical moiety with an azomethine moiety in the presence of a hydrogen atom donor in an intramolecular reaction to form a fused ring system, wherein said azomethine moiety possesses at least one radical stabilizing group, and the azomethine carbon is in the ketone oxidation state or higher.

[0022] As noted above, the process of the present invention is particularly well suited for the synthesis of a wide variety of substituted indoline species. In this regard, the phenyl ring in the indoline ring system may possess one or more heteroatoms and may be fused to one or more aryl or heteroaryl rings, so long as there is an sp^2 hybridizable carbon atom alpha to the point of attachment of the tethered azomethine.

[0023] Thus, in a sixth aspect, there is provided a process for preparing compounds of the formula



wherein each of the groups R^4 , R^5 , and R^8 are independently selected from hydrogen, aryl; heteroaryl; hydrocarbyl; substituted aryl; substituted heteroaryl; substituted hydrocarbyl; heteroatom connected aryl; heteroatom connected hydrocarbyl; heteroatom connected substituted hydrocarbyl; heteroatom connected heteroaryl; heteroatom connected substituted aryl; halo; amino; cyano; hydroxy; carboxy; a group of the formula $-C(O)O-C_1-C_8$ alkyl; a group of the formula $-C(O)R^1$; a group of the formula $-O-R^1$; a group of the formula $-NHR^1$; a group of the formula $-N(R^1)_2$; C_1-C_8 alkoxy; C_1-C_8 alkylthio; or two of R^5 and R^6 , and/or two R^8 groups taken

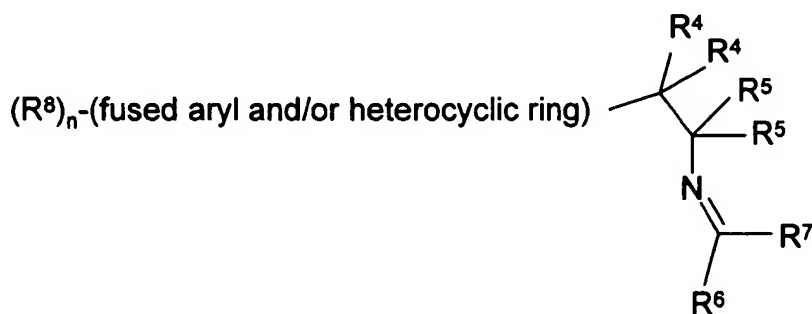
together can form a divalent hydrocarbyl, substituted hydrocarbyl, or be bonded directly to a heteroatom selected from oxygen, nitrogen, or sulfur; and n is from 0 to however many available sites exist on the ring system, for example, 4 in the case of phenyl and 6 in the case of naphthyl;

in addition, two R⁴ groups and/or two R⁵ groups can be taken together represent oxo;

R⁶ and R⁷ are independently selected from aryl, heteroaryl, hydrocarbyl, substituted aryl, substituted heteroaryl, and substituted hydrocarbyl; provided that said groups are bonded via a carbon atom;

and n is from 0 to 4;

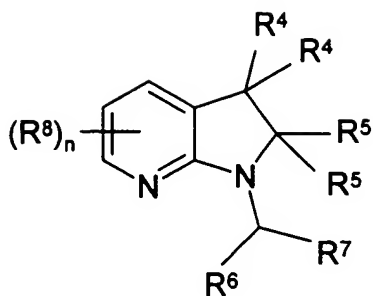
which comprises contacting a compound of the formula



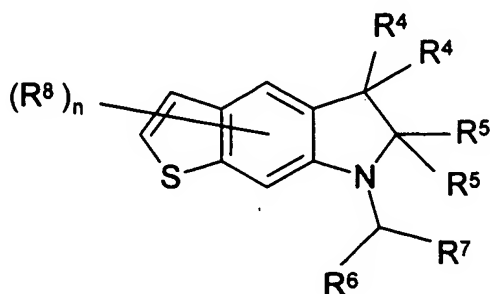
wherein said fused aryl and/or heterocyclic ring possesses a carbon alpha to its point of attachment capable of forming an sp² hybridized carbon radical, said carbon substituted by a group Y, wherein Y is a radical leaving group;

with a free radical initiator in the presence of a hydrogen atom donor.

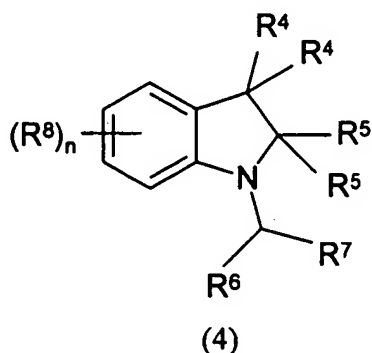
[0024] In this aspect of the invention, it should be appreciated that the groups which are suitable in this process as “fused aryl and/or heterocyclic rings” include aryl and heteroaryl groups as set forth below, optionally substituted by one or more R⁸ groups. By way of example, if the fused aryl and/or heterocyclic ring is pyridine, one possible ring structure which can be obtained via this process has the formula:



[0025] Similarly, if the “fused aryl and/or heterocyclic ring” is benzothiophene, one possible ring system which can be obtained via this process has the formula:



[0026] As noted above, the process of the present invention is especially useful in the preparation of indoline ring systems. Thus, in a fifth aspect, the present invention provides a process for preparing a compound of Formula (4)



wherein each of the groups R^4 , R^5 , and R^8 are independently selected from hydrogen, aryl; heteroaryl; hydrocarbyl; substituted aryl; substituted heteroaryl; substituted hydrocarbyl; heteroatom connected aryl; heteroatom connected hydrocarbyl; heteroatom connected substituted hydrocarbyl; heteroatom connected heteroaryl; heteroatom connected substituted aryl; halo;

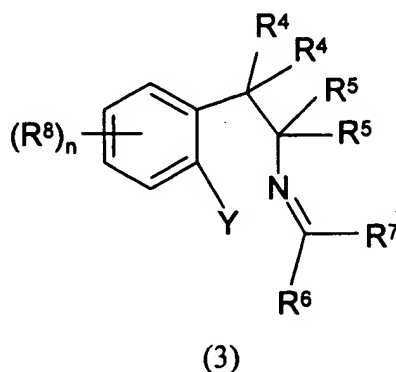
amino; cyano; hydroxy; carboxy; a group of the formula $-C(O)O-C_1-C_8$ alkyl; a group of the formula $-C(O)R^1$; a group of the formula $-O-R^1$; a group of the formula $-NHR^1$; a group of the formula $-N(R^1)_2$; C_1-C_8 alkoxy; C_1-C_8 alkylthio; or two of R^5 and R^6 , or two R^8 groups taken together can form a divalent hydrocarbyl, substituted hydrocarbyl, or be bonded directly to a heteroatom selected from oxygen, nitrogen, or sulfur;

in addition, two R^4 groups and/or two R^5 groups can be taken together represent oxo;

R^6 and R^7 are independently selected from aryl, heteroaryl, hydrocarbyl, substituted aryl, substituted heteroaryl, and substituted hydrocarbyl; provided that said groups are bonded via a carbon atom;

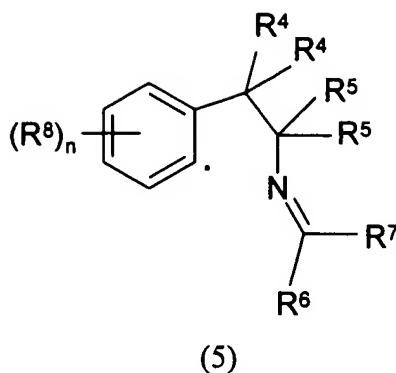
and n is from 0 to 4;

which comprises contacting a compound of Formula (3)



with a free radical initiator in the presence of a hydrogen atom donor,
wherein Y is a radical leaving group.

[0027] In the process of this aspect of the invention, the desired radical may be generated *in situ* using methodology described above with regard to the first aspect. Thus, in a further embodiment, there is provided the free radical intermediate of the Formula (5)



which is useful as an intermediate in the preparation of the compounds of Formula (4).

[0028] In a further preferred embodiment of this aspect, the invention is useful in the preparation of chiral products, *i.e.*, intermediates and products in which the product of the process is enantiomerically enriched. In this regard, in the case where one of the R^5 groups is carboxy, such a chiral synthetic route allows for the preparation of indoline α -amino acids.

[0029] The advantages of the methodology of this invention include the readily-available aryl radical precursors (*e.g.*, aryl halides), readily available phenethyl amine derivatives and carbonyl compounds, as well as the use of radical intermediates and reaction conditions which are pH neutral. Thus, no acid or base addends are necessary to effect the desired transformation. This feature is especially advantageous when the desired indole compound possesses acid or base labile functional groups and/or is comprised of chiral center(s) susceptible to acid/base-catalyzed epimerization. Moreover, there are no electronic restrictions on the substrates (3), and thus acid- and base-sensitive substrates that are electron rich, electron neutral, or electron deficient may be utilized. This addition of aryl radicals to the nitrogen of an azomethine moiety may result in excess of 10:1, more often 20:1 regioselectivity for the desired nitrogen-carbon bond formation. In contrast to other methodologies for amination, the active nitrogen-containing component in each case is an azomethine, and the pi-bond is its reactive feature (as opposed to a sigma N-H bond).

[0030] In the above Formulae (3), (4), and (5), it will be understood that the indeterminate placement of the R^8 groups denotes that there may be none or up to 4 such groups at the carbon atoms of the phenyl ring. Thus, the process of this aspect of the invention should be recognized as a general methodology for the synthesis of indole and substituted indole species. In this regard, the possible substituents from which R^8 , as well as R^4 , R^5 , R^6 , and R^7 may be selected is not particularly limited, so long as such groups do not contain species which have a deleterious

effect on the desired reaction or otherwise serve to quench the sp^2 -hybridized carbon radical prematurely. Insofar as such groups are not limited by acid/base sensitivity or by electronic features, the above process is useful for making a broad range of compounds containing the indole subunit. By way of illustration of the robust nature of the process of the present invention, one R^4 group and one R^5 group could be taken together to form an aziridine ring system.

[0031] Examples of particularly preferred substrates of Formula (3) include homochiral indoline α -amino acids, substituted indolines, *etc.*

[0032] It is this flexibility which renders the above methodologies especially suited for the construction of libraries of compounds, insofar as wide varieties of electron-rich as well as electron-deficient amine and ketone starting materials can be utilized to prepare the azomethine species and in all such cases, provided the above parameters are met, will result in the formation of intramolecular carbon-nitrogen bond formation. Thus, in a further aspect, there is provided a process for the synthesis of a library of compounds having a proline, indoline, or indole subunit, which comprises application of the methodology herein.

[0033] In each of the above aspects of the process of the present invention, the process may be conducted neat or in a nonparticipating solvent such as benzene.

[0034] Also, in each of the above aspects of the invention, the methods may be conducted with the substrate attached to a solid support.

[0035] Other desired reaction conditions for the process of the present invention are not particularly critical and may in any event be chosen and optimized by one of ordinary skill in the art.

[0036] In this disclosure certain chemical groups or compounds are described by certain terms and symbols. These terms are defined as follows:

[0037] Symbols ordinarily used to denote elements in the Periodic Table take their ordinary meaning, unless otherwise specified. Thus, N, O, S, P, and Si stand for nitrogen, oxygen, sulfur, phosphorus, and silicon, respectively.

[0038] A "hydrocarbyl" group means a monovalent or divalent, linear, branched or cyclic group which contains only carbon and hydrogen atoms. Examples of monovalent hydrocarbyls

include the following: C_1 - C_{20} alkyl; C_1 - C_{20} alkyl substituted with one or more groups selected from C_1 - C_{20} alkyl, C_3 - C_8 cycloalkyl, or aryl; C_3 - C_8 cycloalkyl substituted with one or more groups selected from C_1 - C_{20} alkyl, C_3 - C_8 cycloalkyl, or aryl.

[0039] A "substituted hydrocarbyl" refers to a monovalent or divalent hydrocarbyl substituted with one or more heteroatoms. Examples of monovalent substituted hydrocarbyls include: --C(O)R¹³ (wherein R¹³ is hydrocarbyl), --C(O)NR¹³₂ (wherein R¹³ is hydrocarbyl), 2-hydroxyphenyl, 2-methoxyphenyl, 2-ethoxyphenyl, 2-fluorophenyl, 2-chlorophenyl, 2-trifluoromethylphenyl, 2,6-bis(trifluoromethyl)phenyl, 2-(trialkylsiloxy)phenyl, 2-(triarylsiloxy)phenyl, 2,6-bis(diphenylamino)phenyl, 2,6-bis(phenoxy)phenyl, 2-hydroxy-6-phenylphenyl, 2-cyanophenyl, 2-(diphenylamino)phenyl, 4-nitrophenyl, 2-nitrophenyl, --CH₂OR¹³ (wherein R¹³ is hydrocarbyl), cyano, --CH₂NR¹³₂ (wherein R¹³ is hydrocarbyl), and --CH₂OSiR¹³₃ (wherein R¹³ is hydrocarbyl). Also encompassed by the term "substituted hydrocarbyl" are hydrocarbyl groups having one or more groups selected from amide, imido, carbonyl, carboxy, hydroxy, cyano, nitro, halo, alkoxy, alkoxycarbonyl, carboxamido groups, as well as unsubstituted and substituted carboxylic acid ester, unsubstituted and substituted carbamoyl, and substituted imino, as such terms are defined herein. Moreover, such substituted hydrocarbyl groups can form aliphatic ring systems containing one or more heteroatoms and various alkylene linkages such as methylene, ethylene, propylene, etc. Examples of such ring systems include but are not limited to tetrahydrofuran, pyrrolidine, tetrahydropyran, piperidine, and the like.

[0040] Examples of the term "aryl" include a unsubstituted C_6 - C_{14} aryl group as well as a C_6 - C_{14} aryl substituted with one or more groups selected from C_1 - C_{20} alkyl, C_3 - C_8 cycloalkyl, aryl, amide, imido, carbonyl, carboxy, hydroxy, cyano, nitro, halo, alkoxy, alkoxycarbonyl, carboxamido groups, as well as unsubstituted and substituted carboxylic acid ester, unsubstituted and substituted carbamoyl, and substituted imino, as such terms are defined herein, and wherein the term "aryl" preferably denotes a phenyl, naphthyl, or anthracenyl group.

[0041] A "heteroatom" refers to an atom other than carbon or hydrogen. Preferred heteroatoms include oxygen, nitrogen, phosphorus, sulfur, selenium, arsenic, chlorine, bromine, silicon and fluorine. In this regard, the terms "heteroatom connected aryl", "heteroatom connected hydrocarbyl" and like terms indicate that a heteroatom is the point of attachment for such a group. An example of a heteroatom connected hydrocarbyl would thus be butanethiol, an example of a heteroatom connected aryl would be phenoxy, and an example of a heteroatom connected heteroaryl would be 1-pyrrolyl.

[0042] The term "heteroaryl" as used herein preferably refers to heterocyclic aryl rings having one or more heteroatoms. Preferably, such heteroaryl groups are stable 5- to 7-membered monocyclic or bicyclic or 7- to 10-membered bicyclic heterocyclic rings which are saturated partially unsaturated or unsaturated (aromatic), and which consists of carbon atoms and from 1 to 4 heteroatoms independently selected from the group consisting of N, O and S and including any bicyclic group in which any of the above-defined heterocyclic rings is fused to a benzene ring. The nitrogen and sulfur heteroatoms may optionally be oxidized. The heterocyclic ring may be attached to its pendant group at any heteroatom or carbon atom which results in a stable structure. The heterocyclic rings described herein may be substituted on carbon or on a nitrogen atom if the resulting compound is stable. A nitrogen in the heterocycle may optionally be quaternized. It is preferred that when the total number of S and O atoms in the heterocycle exceeds 1, then these heteroatoms are not adjacent to one another. It is preferred that the total number of S and O atoms in the heterocycle is not more than 1. Examples of heterocyclic rings include, but are not limited to, 1H-indazole, 2-pyrrolidonyl, 2H,6H-1,5,2-dithiazinyl, 2H-pyrrolyl, 3H-indolyl, 4-piperidonyl, 4aH-carbazole, 4H-quinoliziny, 6H-1,2,5-thiadiazinyl, acridinyl, azocinyl, benzimidazolyl, benzofuranyl, benzothiofuranyl, benzothiophenyl, benzoxazolyl, benzthiazolyl, benztriazolyl, benztetrazolyl, benzisoxazolyl, benzisothiazolyl, benzimidazolonyl, carbazolyl, 4aH-carbazolyl, .beta.-carbolinyl, chromanyl, chromenyl, cinnolinyl, decahydroquinolinyl, 2H,6H-1,5,2-dithiazinyl, dihydrofuro[2,3-b]tetrahydrofuran, furanyl, furazanyl, imidazolidinyl, imidazoliny, imidazolyl, 1H-indazolyl, indolenyl, indolinyl, indoliziny, indolyl, isobenzofuranyl, isochromanyl, isoindazolyl, isoindolinyl, isoindolyl, isoquinolinyl isothiazolyl, isoxazolyl, morpholinyl, naphthyridinyl, octahydroisoquinolinyl, oxadiazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolidinyl., oxazolyl, oxazolidinylperimidinyl, phenanthridinyl, phenanthrolinyl, phenarsazinyl, phenazinyl, phenothiazinyl, phenoxathiinyl, phenoxazinyl, phthalazinyl, piperazinyl, piperidinyl, pteridinyl, piperidonyl, 4-piperidonyl, pteridinyl, purinyl, pyranyl, pyrazinyl, pyrazolidinyl, pyrazolinyl, pyrazolyl, pyridazinyl, pyridoazole, pyridoimidazole, pyridothiazole, pyridinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolinyl, pyrrolyl, quinazolinyl, quinolinyl, 4H-quinoliziny, quinoxaliny, quinuclidinyl, carbolinyl, tetrahydrofuranyl, tetrahydroisoquinolinyl, tetrahydroquinolinyl, 6H-1,2,5-thiadiazinyl, 1,2,3-thiadiazolyl, 1,2,4-thiadiazolyl, 1,2,5-thiadiazolyl, 1,3,4-thiadiazolyl, thianthrenyl, thiazolyl, thienyl, thienothiazolyl, thienooxazolyl, thienoimidazolyl, thiophenyl, triazinyl, 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,5-triazolyl, 1,3,4-triazolyl, xanthenyl. Preferred heterocycles include, but are not limited to, pyridinyl, furanyl, thienyl, pyrrolyl, pyrazolyl, imidazolyl, indolyl, benzimidazolyl,

1H-indazolyl, oxazolidinyl, benzotriazolyl, benzisoxazolyl, oxindolyl, benzoxazoliny, or isatinoyl.

[0043] The terms "substituted aryl" and "substituted heteroaryl" refers to such aryl and heterocyclic rings substituted by one or more halogen, C₁ -C₆ alkyl, phenoxy, phenyl, hydroxy, amino, C₁ -C₆ alkoxycarbonyl, nitro, C₁ -C₆ alkylsulfonyl, carboxy, cyclohexyl, carbamoyl, cyano, C₁ -C₆ alkylsulfonylamino or C₁ -C₆ alkoxy groups, amido, imido, cyano, nitro, C₁ -C₆ alkoxy, C₁ -C₆ carboxamido groups, as well as unsubstituted and substituted carboxylic acid ester, unsubstituted and substituted carbamoyl, and substituted imino, as such terms are defined herein.

[0044] The term "alkoxycarbonyl" refers to an alkoxy group bonded to a carbonyl function. In other words, the C₂ alkoxycarbonyl group is ethoxycarbonyl. The term "substituted alkoxycarbonyl" refers to a C₁ -C₆ alkoxycarbonyl group substituted with one or more halogen, phenyl, phenoxy, hydroxy, amino, C₁ -C₆ alkoxycarbonyl, carboxy, cyclohexyl, carbamoyl, cyano, C₁ -C₆ alkylsulfonylamino, or C₁ -C₆ alkoxy groups.

[0045] The terms "alkyl" and "alkylene" as used herein preferably refer to C₁ -C₁₂ straight or branched chain alkyl and alkylene groups, respectively. The terms "lower alkenyl" and "lower alkynyl" refer to C₃ -C₆ alkenyl groups and C₃ -C₆ alkynyl groups, respectively.

[0046] The term "unsubstituted and substituted carboxylic acid ester" refers to a C₁ -C₈ alkyl, C₃ -C₈ cycloalkyl or aryl oxycarbonyl group, preferably containing from 2 to 10 carbon atoms and optionally substituted with halogen, C₁ -C₆ alkoxy, C₃ -C₈ cycloalkyl, aryl, aryloxy, C₁ -C₆ alkyl, cyano, C₁ -C₆ alkanoyloxy, hydroxy or C₁ -C₆ alkoxycarbonyl.

[0047] The term "unsubstituted and substituted carbamoyl" refers to an alkyl (or substituted alkyl) amino carbonyl group, preferably containing from 2 to 10 carbon atoms.

[0048] The term "substituted imino" refers to an imino group substituted with a group selected from hydrogen, C₁ -C₆ alkyl, C₃ -C₈ cycloalkyl or aryl.

[0049] The term "radical leaving group" will be understood by those skilled in the art to denote groups such as iodo, bromo, diazonium salts, and the like.

[0050] The term "solid support" as used herein preferably refers to solid supports known in the art of synthetic chemistry as inert materials which are attached to a substrate, upon which one or more synthetic manipulations is to be carried out. This material upon which the processes of the invention are performed are referred to as solid supports, beads and resins. These terms are intended to include: beads, pellets, disks, fibers, gels or particles such as cellulose beads, pore-glass beads, silica gels, polystyrene beads optionally cross-linked with divinylbenzene and optionally grafted with polyethylene glycol and optionally functionalized with amino, hydroxy, carboxy or halo groups, grafted co-poly beads, polyacrylamide beads, latex beads, dimethylacrylamide beads optionally cross-linked with N,N'-bis-acryloyl ethylene diamine, glass particles coated with hydrophobic polymer, *etc.*, *i.e.*, material having a rigid or semi-rigid surface and soluble supports such as low molecular weight non-cross-linked polystyrene.

[0051] The following examples are set forth merely as illustrations of the invention and are not intended to be considered limiting as to the scope thereof.

Experimental Section

[0052] Flame-dried (under vacuum) glassware was used for all non-aqueous reactions. All reagents and solvents were commercial grade and purified prior to use when necessary. Diethyl ether (Et₂O), tetrahydrofuran (THF), dichloromethane (CH₂Cl₂), and benzene (C₆H₆) were dried by passage through a column of activated alumina as described by Grubbs (See Pangborn, A.B.; Giardello, M.A.; Grubbs, R.H.; Rosen, R.K.; Timmers, F.J. *Organometallics* **1996**, *15*, 1518–1520.) Benzene was additionally passed through a column containing activated Q-5 reactant. Solvents other than benzene were degassed using the freeze-pump-thaw method when necessary. All additional solvents were dried by distillation from calcium hydride when necessary. Molecular sieves (spheres, 4Å) were calcined at 400 °C and stored at room temperature in an air-tight container. AIBN was recrystallized prior to use, and tri-*n*-butyl tin hydride (ⁿBu₃SnH) was used as received from Aldrich. Preparations for previously unreported phenethyl amine derivatives used in this study will be reported later in an Article after optimization.

[0053] Thin layer chromatography (TLC) was performed using glass-backed silica gel (250 μ) plates and flash chromatography utilized 230–400 mesh silica gel from Scientific Adsorbents. Neutral Alumina was used as received from Scientific Adsorbents for chromatography of acid-sensitive intermediates or products. Products were visualized by UV light, iodine, and/or the use

of ceric ammonium molybdate, potassium permanganate, ninhydrin, *p*-anisaldehyde, and potassium iodoplatinate solutions.

[0054] IR spectra were recorded on a Nicolet Avatar 360 spectrophotometer. Liquids and oils were analyzed as neat films on a salt plate (transmission), whereas solids were applied to a diamond plate (ATR). Nuclear magnetic resonance spectra (NMR) were acquired on either a Varian Inova-400 or VXR-400 instrument. Chemical shifts are measured relative to tetramethylsilane, as judged by the residual partially deuterated solvent peak. Mass spectra were obtained using a Kratos MS-80 spectrometer using the ionization technique indicated. Combustion analyses were performed on a Perkin-Elmer 2400 Series II CHNS/O Analyzer.

[0055] Ratios of diastereomers and isomeric products were measured directly from integration of ^1H NMR absorptions of protons common to the components. Precision was checked by varying the relaxation delay for measurements on the same compound. Where possible, ratios were corroborated using GC-mass spectrometry. Peak assignments were made from authentic samples in every case. Ratios reported generally represent a lower limit defined by multiple runs.

General Procedure for Ketimine Condensations

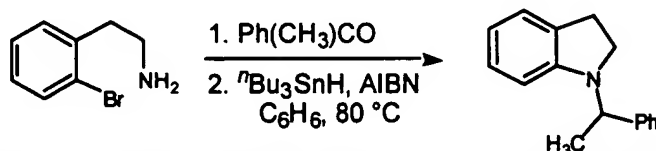
[0056] A rapidly stirred benzene solution of the amine (0.5 M), ketone (0.5 M), and 4 Å MS (1:1 w/w) was stirred at 25 °C until complete conversion was achieved, as evidenced by ^1H NMR. The mixture was filtered through a pad of Celite and washed with Et₂O or benzene. The solvent was removed *in vacuo* to give the analytically pure ketimine which was used immediately.

[0057] The same procedure was used when the benzophenone ketimine was desired, except benzophenone imine (Pickard, P.L.; Tolbert, T.L., in "Organic Syntheses"; Wiley: NY, 1973, Collective Vol. 5, pp. 520-2) was used in place of the ketone. (O'Donnell, M.J.; Polt, R.L. *J. Org. Chem.* **1982**, *47*, 2663.)

General Procedure for Aryl Aminations

[0058] A benzene solution of the ketimine (0.01 M) was warmed to 85 °C in a round-bottomed flask equipped with a condenser. A benzene solution (1 mL) of $n\text{Bu}_3\text{SnH}$ (1.1 equiv) and AIBN (0.4 equiv) was loaded into a gas-tight syringe and was attached to a syringe pump. The syringe needle was attached through a septum at the top of the condenser (w/ N_2 line) so that the solution droplets would fall directly into the refluxing benzene. Following the addition, the reaction mixture was refluxed for an additional period (~1 h) and cooled to room temperature. At this point, an aliquot was removed, concentrated, and component ratios were measured by ^1H NMR and/or GC-MS. The solution was treated with NaBH_4 (1.1 equiv) and the slurry was stirred 4–5 hours. The mixture was concentrated *in vacuo*, diluted with Et_2O , and washed with water. The organic layer was separated, dried (MgSO_4), and concentrated to furnish an oil. Flash chromatography of the crude mixture provided the analytically pure targeted compounds.

Example 1

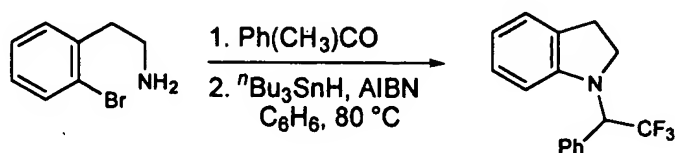


[0059] *N*-(1-Methylbenzyl)indoline (2a). Following the general procedure, *o*-bromophenethylamine (46 mg, 231 mmol), acetophenone (27 μL , 231 mmol), and 4Å MS were stirred in benzene (1 mL) at room temperature for 12 h. Filtering of the mixture through Celite and removal of the solvent provided the ketimine as a >95:5 mixture of stereoisomers. IR (film) 3056, 1633 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.78 (m, 2H), 7.57 (d, J = 6.9 Hz, 2H), 7.40 (t, J = 3.1 Hz, 3H), 7.34 (d, J = 6.0 Hz, 1H), 7.26 (t, J = 6.3 Hz, 1H), 7.10 (t, J = 6.0 Hz, 1H), 3.79 (dd, J = 7.8, 7.8 Hz, 2H), 3.22 (dd, J = 7.8, 7.8 Hz, 2H), 2.17 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 166.0, 142.0, 140.0, 133.0, 131.7, 129.7, 128.5, 128.1, 127.6, 126.9, 52.2, 37.8, 18.1; HRMS (EI): Exact mass calcd for $\text{C}_{16}\text{H}_{16}\text{BrN}$ $[\text{M}+\text{H}]^+$, 302.0544. Found 302.0516.

[0060] A three hour addition of a $n\text{Bu}_3\text{SnH}$ (68 μL , 0.25 mmol) and AIBN (15 mg, 93 μmol) solution in benzene (0.7 mL) to a refluxing solution of the unpurified ketimine (69.6 mg, 231 μmol) in benzene (23 mL) delivered, after flash chromatography (2% CH_2Cl_2 in hexanes), 44.9 mg (87%) of the desired indoline as a colorless oil. R_f = 0.62 (30% CH_2Cl_2 /hexanes); IR (film)

3046, 1606 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.41 (d, $J = 7.5$ Hz, 2H), 7.29 (t, $J = 7.2$ Hz, 2H), 7.26 (t, $J = 7.2$ Hz, 1H), 7.08 (d, $J = 7.2$ Hz, 1H), 7.00 (t, $J = 7.8$ Hz, 1H), 6.6 (t, $J = 7.2$ Hz, 1H), 6.36 (d, $J = 7.8$ Hz, 1H), 4.68 (q, $J = 6.8$ Hz, 1H), 3.42 (ddd, $J = 18.1, 9.1, 0$ Hz, 1H), 3.35 (ddd, $J = 15.7, 7.4, 0$ Hz, 1H), 2.96 (dd, $J = 8.5, 8.3$ Hz, 2H), 2.58 (d, $J = 6.8$ Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 151.1 143.0 128.7, 127.4, 127.3, 127.1, 124.6, 117.2, 107.5, 54.8, 48.2, 28.5, 16.8; HRMS (EI): Exact mass calcd for $\text{C}_{16}\text{H}_{17}\text{N}$ $[\text{M}]^+$, 223.1361. Found 223.1366.

Example 2

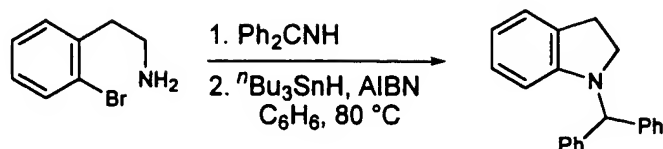


[0061] N-(1-Trifluoromethyl)benzyl indoline (2b). Following the general procedure, o-bromophenethylamine (852 mg, 4.25 mmol), trifluoroacetophenone (590 μL , 4.21 mmol), and 4Å MS were stirred in toluene (10 mL) at room temperature for 12 h to provide the ketimine as a >95:5 mixture of stereoisomers. IR (film) 3062, 1669 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.49 (d, $J = 7.9$ Hz, 1H), 7.44 (t, $J = 7.4$ Hz, 1H), 7.38 (t, $J = 7.1$ Hz, 2H), 7.24 (t, $J = 7.4$ Hz, 1H), 7.18 (d, $J = 6.0$ Hz, 1H), 7.10 (t, $J = 6.0$ Hz, 1H), 6.94 (d, $J = 7.1$ Hz, 2H), 3.69 (dd, $J = 7.1, 7.1$ Hz, 2H), 3.13 (dd, $J = 7.1, 7.1$ Hz, 2H); ^{13}C NMR (100 MHz, CDCl_3) ppm 138.4, 133.0, 131.7, 130.3, 130.1, 128.8, 128.4, 127.7, 127.5, 124.9, 52.8 36.8; HRMS (EI): Exact mass calcd for $\text{C}_{16}\text{H}_{13}\text{F}_3\text{N}$ $[\text{M}-\text{Br}]^+$, 276.1000. Found 276.1003.

[0062] A three hour addition of $^t\text{Bu}_3\text{SnH}$ (184 μL , 682 μmol) and AIBN (41 mg, 248 μmol) solution in benzene (2.5 mL) to a refluxing solution of the unpurified ketimine (220 mg, 620 μmol) in benzene (62 mL) delivered, after flash chromatography (5% CH_2Cl_2 in hexanes), 133 mg (77%) of the desired indoline as a colorless oil. $R_f = 0.35$ (5% EtOAc/hexanes); IR (film) 3031, 1607 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.40 (m, 5H), 7.11 (t, $J = 7.9$ Hz, 1H), 7.09 (d, $J = 7.4$ Hz, 1H), 6.71 (t, $J = 7.4$ Hz, 1H), 6.62 (d, $J = 7.9$ Hz, 1H), 5.24 (q, $J = 8.7$ Hz, 1H), 3.62 (dd, $J = 14.2, 8.7$, Hz, 1H), 3.19 (dd, $J = 10.1, 8.9$, Hz, 1H), 3.01 (m, 2H); ^{13}C NMR (100 MHz, CDCl_3) ppm 150.5, 131.9, 129.3, 129.0, 127.6, 127.5, 125.1, 124.7, 118.5, 106.4, 62.0 (q, $J =$

30.5 Hz, 1C), 48.6, 28.5; HRMS (EI): Exact mass calcd for $C_{16}H_{14}F_3N [M]^+$, 277.1078.
Found 277.1071.

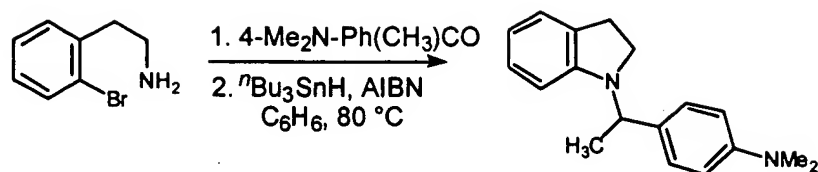
Example 3



[0063] ***N*-(1-Phenylbenzyl)indoline (2c).** Following the general procedure, *o*-bromophenethylamine (471 mg, 2.35 mmol) and benzophenone imine (426 mg, 2.35 mmol) were stirred for 12 h in dichloromethane (4 mL). Removal of the solvent provided the ketimine. IR (film) 3056, 1660, 1623 cm^{-1} ; 1H NMR (400 MHz, $CDCl_3$) δ 7.61 (d, $J = 6.8$ Hz, 2H), 7.49 (d, $J = 7.0$ Hz, 1H), 7.40 (m, $J = 3.8$ Hz, 3H), 7.35 (m, $J = 7.0$ Hz, 3H), 7.24 (t, $J = 5.8$ Hz, 1H), 7.19 (d, 6.8 Hz, 1H), 7.05 (t, $J = 5.8$ Hz, 1H), 6.98 (m, $J = 3.8$ Hz, 2H), 3.68 (dd, $J = 7.2, 7.2$ Hz, 2H), 3.14 (dd, $J = 7.4, 7.4$ Hz, 2H); ^{13}C NMR (100 MHz, $CDCl_3$) ppm 169.0, 140.0, 139.8, 136.9, 132.9, 131.6, 130.1, 128.65, 128.61, 128.4, 128.2, 127.9, 127.4, 125.0, 53.6, 37.9; HRMS (CI): Exact mass calcd for $C_{21}H_{18}BrN [M+H]^+$, 364.0701. Found 364.0596.

[0064] A three hour addition of a nBu_3SnH (70 μL , 254 μmol) and AIBN (5 mg, 9.2 μmol) solution in benzene (1.0 mL) to a refluxing solution of the unpurified ketimine in benzene (23 mL) delivered, after flash chromatography (2% CH_2Cl_2 in hexanes), 56.8 mg (86%) of the desired indoline as a white solid. mp 62–63 $^\circ C$. $R_f = 0.53$ (10% EtOAc in hexanes); IR (film) 3025, 1605 cm^{-1} ; 1H NMR (400 MHz, $CDCl_3$) δ 7.35 (m, 10H), 7.09 (d, $J = 7.0$ Hz, 1H), 6.92 (t, $J = 7.5$ Hz, 1H), 6.64 (t, $J = 7.0$ Hz, 1H), 6.20 (d, $J = 7.5$ Hz, 1H), 5.55 (s, 1H), 3.20 (dd, $J = 8.3, 8.2$ Hz, 2H), 2.96 (dd, $J = 8.3, 8.2$ Hz, 2H), ^{13}C NMR (100 MHz, $CDCl_3$) ppm 152.1, 141.5, 130.5, 128.7, 128.6, 127.4, 127.3, 124.5, 117.7, 66.8, 51.6, 28.5; HRMS (EI): Exact mass calcd for $C_{21}H_{19}N [M]^+$, 285.1517. Found 285.1520.

Example 4

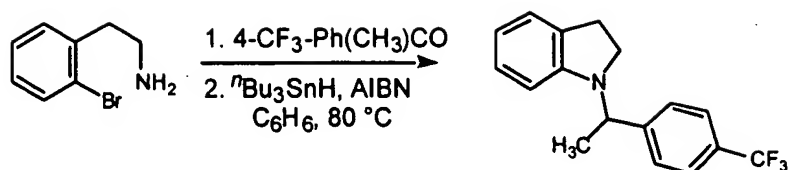


[0065] *N*-[(1-Methyl)-4-dimethylaminobenzyl]-indoline (2d). According to the general procedure, *o*-bromophenethylamine (100 mg, 500 μ mol), aceto(*p*-*N,N'*-dimethylamino)phenone (81.5 mg, 500 μ mol), and 4Å MS were stirred in benzene (6 mL) at room temperature for 4 h to provide the ketimine as a >95:5 mixture of stereoisomers. IR (film) 3050, 1602 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.63 (d, J = 9.0 Hz, 2H), 7.46 (dd, J = 8.1, 1.1 Hz, 1H), 7.25 (dd, J = 7.6, 1.5 Hz, 1H), 7.15 (dt, J = 7.4, 1.1 Hz, 1H), 6.98 (dt, J = 6.7, 1.6 Hz, 1H), 6.61 (d, J = 9.0 Hz, 2H), 3.65 (t, J = 7.4 Hz, 2H), 3.10 (t, J = 7.4 Hz, 2H), 2.91 (s, 6H), 2.03 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 165.4, 151.6, 143.1, 140.3, 132.9, 131.6, 129.5, 128.0, 124.9, 111.7, 51.9, 40.6, 38.01, 15.09; HRMS (EI): Exact mass calcd for $\text{C}_{18}\text{H}_{21}\text{BrN}_2$ $[\text{M}]^+$, 334.0888. Found 344.0824.

[0066] A five hour addition of a $n\text{Bu}_3\text{SnH}$ (56 μL , 207 μmol) and AIBN (12 mg, 75 μmol) solution in benzene (1 mL) to a refluxing solution of the unpurified ketimine (65 mg, 188 μmol) in benzene (18 mL) delivered, after flash chromatography (50% CH_2Cl_2 in hexanes), 45 mg (90%) of the desired indoline as a crystalline solid, mp 79-82 $^\circ\text{C}$; R_f = 0.15 (50%

CH_2Cl_2 /hexanes; IR (film) 2960, 1653 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.28 (t, J = 4.2 Hz, 2H), 7.03 (dd, J = 17.5, 7.2 Hz, 2H), 6.73 (d, J = 8.8 Hz, 2H), 6.50 (t, J = 7.3 Hz, 1H), 6.44 (d, J = 7.9 Hz, 1H), 4.71 (q, J = 6.8 Hz, 1H), 3.37 (q, J = 18.0, 9.2 Hz, 1H), 3.27 (q, J = 15.2, 8.4 Hz, 1H), 2.95 (s, 6H), 2.93 (t, J = 8.0 Hz, 2H), 1.51 (d, J = 6.9 Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 128.2, 127.4, 124.8, 116.8, 112.6, 107.3, 92.1, 86.3, 53.9, 47.9, 41.0, 28.4, 16.3; HRMS (EI): Exact mass calcd for $\text{C}_{18}\text{H}_{22}\text{N}_2$ $[\text{M}]^+$, 266.1783. Found 266.1774.

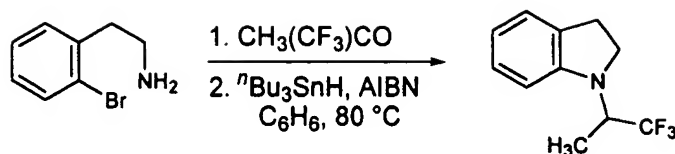
Example 5



[0067] *N*-[(1-Methyl)-4-dimethylaminobenzyl]-indoline (2e). Following the general procedure, *o*-bromophenethylamine (429 mg, 2.14 mmol), *p*-trifluoromethyl acetophenone (404 mg, 2.14 mmol), and 4Å MS were stirred in benzene (5 mL) at room temperature for 4.5 h to provide the ketimine as a 93:7 mixture of stereoisomers. IR (film) 3064, 1636 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.87 (d, J = 8.2 Hz, 2H), 7.46 (d, J = 8.3 Hz, 2H), 7.57 (d, J = 7.8 Hz, 1H), 7.38 (s, 1H), 7.33-7.21 (m, 1H), 7.10 (t, J = 7.7 Hz, 1H), 3.81 (t, J = 7.3 Hz, 2H), 3.23 (t, J = 7.3 Hz, 2H); ^{13}C NMR (100 MHz, CDCl_3) ppm 164.9, 144.6, 139.8, 133.0, 131.7, 128.6, 128.7, 129.6, 127.6, 127.2, 125.6, 124.9, 52.3, 37.7, 15.7; HRMS (FAB): Exact mass calcd for $\text{C}_{17}\text{H}_{16}\text{BrF}_3\text{N}$ $[\text{M}+\text{H}]^+$, 370.0418. Found 370.0428.

[0068] A three hour addition of a $n\text{-Bu}_3\text{SnH}$ (91 μL , 337 μmol) and AIBN (20 mg, 123 μmol) solution in benzene (1 mL) to a refluxing solution of the unpurified ketimine (113 mg, 307 μmol) in benzene (31 mL) delivered, after flash chromatography (20% CH_2Cl_2 in hexanes), 65 mg (72%) of the desired indoline as a colorless oil; R_f = 0.40 (10% EtOAc/hexanes; IR (film) 3048, 1606 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.61 (d, J = 8.3 Hz, 2H), 7.54 (d, J = 8.3 Hz, 2H), 7.10 (d, J = 6.9 Hz, 1H), 7.01 (t, J = 7.5 Hz, 1H), 6.65 (t, J = 7.1 Hz, 1H), 6.31 (d, J = 7.9 Hz, 1H), 4.73 (q, J = 6.8 Hz, 1H), 3.43 (dd, J = 17.9, 8.7 Hz, 1H), 3.37 (dd, J = 15.7, 7.4 Hz, 1H), 2.99 (t, J = 8.3 Hz, 2H), 1.57 (d, J = 6.9 Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 151.3, 147.5, 130.4, 127.5, 127.4, 125.7, 125.6, 124.8, 54.9, 48.5, 28.5, 17.1; HRMS (EI): Exact mass calcd for $\text{C}_{17}\text{H}_{16}\text{F}_3\text{N}$ $[\text{M}]^+$, 291.1235. Found 291.1234.

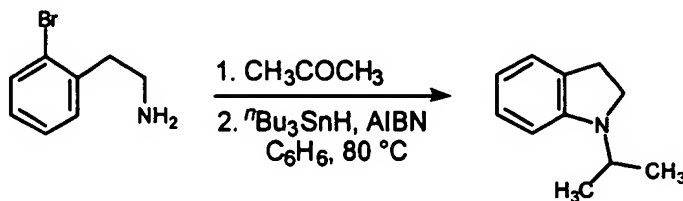
Example 6



[0069] *N*-(1-Trifluoroethyl)-indoline (2f). Following the general procedure, *o*-bromophenethylamine (200 mg, 999 μmol), trifluoroacetone (168 mg, 1.5 mmol), and 4Å MS were stirred in benzene (5 mL) at room temperature for 4 h to provide the ketimine as a >95:5 mixture of stereoisomers. IR (film) 3059, 1686 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.57 (d, $J = 8.3$ Hz, 1H), 7.29–7.22 (m, 2H), 7.13 (dt, $J = 6.8, 2.4$ Hz, 1H), 3.75 (t, $J = 7.3$ Hz, 2H), 3.17 (t, $J = 7.3$ Hz, 2H); 1.85 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 157.3 (q, $J = 33.6$ Hz), 138.6, 133.1, 131.8, 127.8, 124.7, 121.4, 118.6, 51.4, 36.6, 12.6; HRMS (EI): Exact mass calcd for $\text{C}_{11}\text{H}_{13}\text{BrF}_3\text{N}$ $[\text{M}+\text{H}]^+$, 296.0086. Found 296.0088.

[0070] A three hour addition of a $n\text{Bu}_3\text{SnH}$ (133 μL , 495 μmol) and AIBN (30 mg, 180 μmol) solution in benzene (1 mL) to a refluxing solution of the unpurified ketimine (132 mg, 450 μmol) in benzene (44 mL) provided, after flash chromatography (100% hexanes), 80 mg (83%) of the desired indoline as a colorless oil. $R_f = 0.15$ (hexanes); IR (film) 3050, 1490 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.07 (t, $J = 7.9$ Hz, 2H), 6.65 (t, $J = 7.3$ Hz, 1H), 6.44 (d, $J = 7.8$ Hz, 1H), 4.14 (m, 1H), 3.54 (t, $J = 8.5$ Hz, 2H), 3.03 (t, $J = 8.6$ Hz, 2H), 1.38 (d, $J = 7.0$ Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 150.3, 129.3, 127.5, 125.0, 118.2, 114.0, 106.3, 52.8 (q, $J = 28.2$ Hz), 47.1, 28.5, 10.5; HRMS (EI): Exact mass calcd for $\text{C}_{11}\text{H}_{12}\text{F}_3\text{N}$ $[\text{M}]^+$, 215.0922. Found 215.0920.

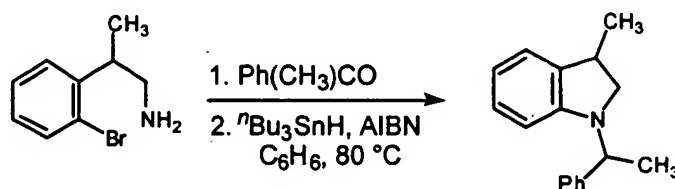
Example 7



[0071] ***N*-(*iso*-Propyl)-indoline (2g).** Following the general procedure, *o*-bromophenethylamine (150 mg, 750 μ mol), acetone (44 mg, 750 μ mol) and 4Å MS were stirred in benzene (5 mL) at room temperature for 4 h to provide the ketimine. IR (film) 3055, 1653 cm^{-1} ^1H NMR (400 MHz, CDCl_3) δ 7.50 (d, J = 8.3 Hz, 1H), 7.24–7.18 (m, 2H), 7.04 (td, J = 6.9, 2.3 Hz, 1H), 3.46 (t, J = 7.5 Hz, 2H), 3.06 (t, J = 7.7 Hz, 2H), 1.99 (s, 3H), 1.71 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 168.3, 139.9, 133.0, 131.4, 128.1, 127.6, 124.9, 51.6, 37.7, 29.5, 18.6; HRMS (EI) Calcd for $\text{C}_{11}\text{H}_{15}\text{BrN}$ $[\text{MH}]^+$, 240.0310. Found 240.0389

[0072] A three-hour addition of $^n\text{Bu}_3\text{SnH}$ (154 μL , 573 μmol) and AIBN (34 mg, 208 μmol) solution in benzene (1 mL) to a refluxing solution of the unpurified ketimine (125 mg, 521 μmol) in benzene (50 mL) delivered, after flash chromatography (30% CH_2Cl_2 /Hexanes) 0.024 g (30%) of the desired indoline as a colorless oil. R_f = 0.1 (30% CH_2Cl_2 /Hexanes); IR (film) 3047, 1607 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.04 (t, J = 6.7 Hz, 2H), 6.59 (t, J = 7.4 Hz, 1H), 6.42 (d, J = 8.1 Hz, 1H), 3.83 (sep, J = 6.6 Hz, 1H), 3.33 (t, J = 8.5 Hz, 2H), 2.93 (t, J = 8.3 Hz, 2H), 1.15 (d, J = 6.7 Hz, 6H); ^{13}C NMR (100 MHz, CDCl_3) ppm 151.5, 130.5, 127.5, 124.6, 117.1, 107.3, 46.0, 45.7, 28.4, 18.4; HRMS (EI) Exact mass calcd for $\text{C}_{11}\text{H}_{15}\text{N}$ $[\text{M}]^+$, 161.1204. Found 161.1201.

Example 8

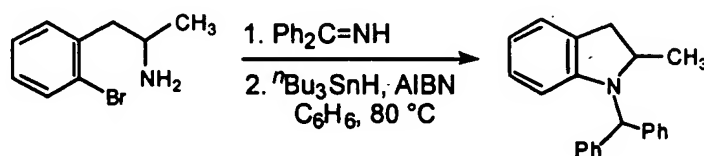


[0073] ***N*-(1-Methylbenzyl)-3-methylindoline (5a).** Following the general procedure, 2-(*o*-bromophenyl)-2-methyl-ethylamine (353 mg, 1.65 mmol), acetophenone (192 μL , 1.64 mmol), and 4Å MS were stirred in toluene (5 mL) at room temperature for 12 h to provide the ketimine as a >95:5 mixture of stereoisomers. IR (film) 3057, 1633 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.80 (m, 2H), 7.59 (d, J = 6.9 Hz, 1H), 7.39 (m, 4H), 7.31 (t, J = 6.9 Hz, 1H), 7.08 (t, J = 6.1 Hz, 1H), 3.76 (dd, J = 10.5, 6.3, Hz, 2H), 3.57 (dq, J = 13.8, 6.9 Hz, 1H), 2.19 (s, 3H), 1.45 (d, J = 6.7 Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 165.5, 144.9, 141.4, 133.0, 129.6, 128.6, 128.4, 128.3, 127.8, 127.6, 126.8, 125.1, 58.1, 39.9, 18.8, 15.7; HRMS (EI): Exact mass calcd for $\text{C}_{17}\text{H}_{19}\text{BrN}$ $[\text{M}+\text{H}]^+$, 316.0701. Found 316.0588.

[0074] A three hour addition of $n\text{Bu}_3\text{SnH}$ (84 μL , 314 μmol) and AIBN (18.7 mg, 114 μmol) benzene solution (1 mL) to a refluxing solution of the unpurified ketimine (90.3 mg, 286 μmol) in benzene (28 mL) afforded, after flash chromatography (1% ether in hexanes), 58.4 mg (86%) of the desired indoline as a yellow oil (58:42 mixture of diastereomers). R_f = 0.6 (10%

EtOAc/hexanes); IR (film) 3024, 1605 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.43 (d, J = 7.5 Hz, 4H), 7.35 (t, J = 7.5 Hz, 6H), 7.03 (d, J = 8.0 Hz, 2H), 7.01 (t, J = 4.3 Hz, 2H), 6.65 (t, J = 7.4 Hz, 2H), 6.36 (d, J = 4.3 Hz, 2H), 4.74 (q, J = 6.9 Hz, 1H), 4.68 (q, J = 6.8 Hz, 1H), 3.62 (dd, J = 8.8, 8.8 Hz, 1H), 3.45 (dd, J = 8.6, 8.6 Hz, 1H), 3.30 (dq, J = 14.6, 7.3 Hz, 1H), 2.95 (dd, J = 8.8, 8.8 Hz, 2H), 1.57 (d, J = 7.0 Hz, 3H), 1.52 (d, J = 6.9 Hz, 3H), 1.33 (d, J = 6.7 Hz, 3H), 1.27 (d, J = 6.8 Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 151.5, 150.8, 146.4, 143.5, 142.8, 135.4, 135.3, 128.68, 128.62, 127.57, 127.53, 127.2, 127.1, 123.5, 123.1, 117.3, 117.0, 107.5, 107.2, 56.5, 56.0, 54.9, 54.3, 35.0, 19.9, 18.3, 17.2, 16.5; HRMS (EI): Exact mass calcd for $\text{C}_{17}\text{H}_{19}\text{N}$ $[\text{M}]^+$, 237.1517. Found 237.1509.

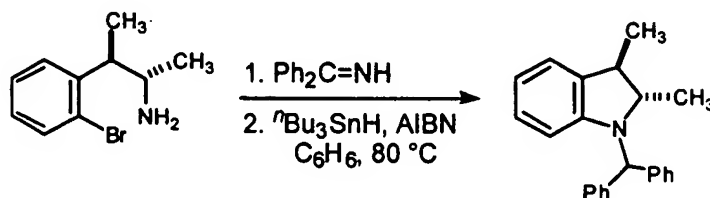
Example 9



[0075] *N*-(1-Phenylbenzyl)-2-methylindoline (5b). Following the general procedure, 2-(*o*-bromophenyl)-1-methylethylamine (50 mg, 234 μmol), benzophenone imine (42 mg, 233 μmol) and 4Å MS were stirred in dichloromethane (2.5 mL) at room temperature for 3 days under argon atmosphere to provide the ketimine as a solid (mp 59–61 $^\circ\text{C}$); IR (film) 3057, 1726, 1661 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.59 (d, J = 6.9 Hz, 2H), 7.45 (d, J = 7.4 Hz, 1H), 7.37–7.30 (m, 6H), 7.20–7.14 (m, 2H), 7.03 (dt, J = 6.9, 1.4 Hz, 1H), 6.63 (d, J = 6.7 Hz, 2H), 3.81 (qd, J = 7.8, 6.2, Hz, 1H), 3.03 (d, J = 4.5 Hz, 1H), 3.01 (d, J = 7.1 Hz, 1H), 1.30 (d, J = 6.2 Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 167.1, 140.2, 139.4, 137.3, 132.6, 130.3, 129.9, 128.6, 128.4, 128.2, 128.1, 127.8, 127.6, 127.1, 125.4, 57.4, 44.8, 22.4; HRMS (EI): Exact mass calcd for $\text{C}_{22}\text{H}_{20}\text{BrN}$ $[\text{M}]^+$, 377.0779. Found 377.0630.

[0076] To a refluxing solution of the unpurified ketimine (20 mg, 53 μmol) in benzene (9 mL) was added $n\text{Bu}_3\text{SnH}$ (15.6 μL , 58.2 μmol), followed by a four-hour addition of AIBN (10.4 mg, 63.5 μmol) dissolved in benzene (2 mL). After the first 2 h of the reaction, another lot of $n\text{Bu}_3\text{SnH}$ (15.6 μL , 58.2 μmol) was added to the reaction mixture. After the complete addition of AIBN, the reaction mixture was refluxed for 1 h, followed by removal of solvent. Flash chromatography (5% CH_2Cl_2 in hexanes) yielded the desired indoline (13.1 mg, 78%) as a solid (mp 58–61 $^\circ\text{C}$); IR (film) 3060, 1635 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.41 (d, $J = 7.6$ Hz, 2H), 7.37–7.26 (m, 8H), 7.04 (d, $J = 7.1$ Hz, 1H), 6.82 (t, $J = 7.5$ Hz, 1H), 6.59 (t, $J = 7.3$ Hz, 1H), 5.97 (d, $J = 7.9$ Hz, 1H), 5.65 (s, 1H), 3.83 (qd, $J = 15.0, 6.2$ Hz, 1H), 3.25 (dd, $J = 15.6, 9.0$ Hz, 1H), 2.65 (dd, $J = 15.4, 6.9$ Hz, 1H), 1.16 (d, $J = 6.0$ Hz, 3H); ^{13}C NMR (75 MHz, CDCl_3) ppm 138.9, 129.2, 129.0, 128.8, 127.6, 127.2, 124.7, 117.4, 109.4, 100.4, 65.5, 52.0, 37.6, 30.1, 21.0; HRMS (EI): Exact mass calcd for $\text{C}_{22}\text{H}_{21}\text{N}$ $[\text{M}]^+$, 299.1674. Found 299.1682.

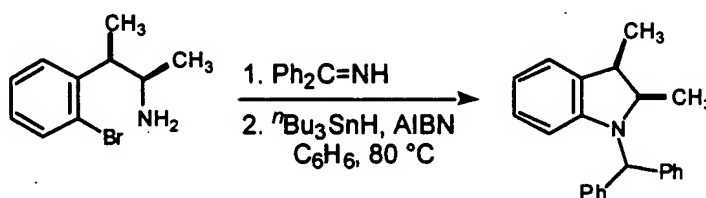
Example 10



[0077] *N*-(1-Phenylbenzyl)-2,3-*trans*-dimethylindoline (5c). Following the general procedure, 2-(*o*-bromophenyl)-1,2-*cis*-dimethylethylamine (127 mg, 562 μmol), benzophenone imine (102 mg, 562 μmol), and 4Å MS were stirred in dichloromethane (1.5 mL) at room temperature for 7.5 days under argon atmosphere to provide the ketimine as an oil; IR (film) 3079, 1661 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.52–7.48 (m, 3H), 7.44–7.40 (m, 3H), 7.34–7.27 (m, 3H), 7.19 (q, $J = 7$ Hz, 1H), 7.18–7.16 (m, 1H), 7.00–6.95 (m, 3H), 3.64 (q, $J = 6.3$ Hz, 2H), 3.62 (q, $J = 6.3$ Hz, 1H), 1.44 (d, $J = 6.6$ Hz, 3H), 1.41 (d, $J = 6.2$ Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 144.6, 140.6, 137.3, 132.7, 129.8, 129.6, 128.7, 128.4, 128.1, 127.5, 62.4, 53.6, 44.8, 19.7, 17.7; HRMS (EI): Exact mass calcd for $\text{C}_{23}\text{H}_{23}\text{BrN}$ $[\text{M}+\text{H}]^+$, 392.1016. Found 392.0913.

[0078] To a refluxing solution of the unpurified ketimine (12.6 mg, 32 μ mol) in benzene (5.5 mL) was added n Bu₃SnH (9.5 μ L, 35 μ mol), followed by a 6 h addition of AIBN (13 mg, 80 μ mol) dissolved in benzene (1 mL). After the first 3 h of the reaction, another aliquot of n Bu₃SnH (9.5 μ L, 35.4 μ mol) was added. After complete addition of AIBN, the reaction mixture was refluxed for further 1 h followed by evaporation of solvent under vacuum. Flash chromatography (5% CH₂Cl₂ in hexanes) yielded the desired indoline (7.9 mg, 79%) as a gum; IR (film) 2958, 1684 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.39 (d, J = 7.2 Hz, 2H), 7.35-7.26 (m, 8H), 7.02 (d, J = 7.2 Hz, 1H), 6.83 (t, J = 7.6 Hz, 1H), 6.61 (t, J = 7.3 Hz, 1H), 5.99 (d, J = 7.9 Hz, 1H), 5.69 (s, 1H), 3.28 (dq, J = 6.3, 6.3 Hz, 1H), 2.87 (dq, J = 6.9, 6.9 Hz, 1H), 1.27 (d, J = 6.4 Hz, 3H), 1.19 (d, J = 6.2 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) ppm 150.4, 141.1, 140.5, 134.3, 128.9, 128.6, 127.4, 123.2, 117.2, 109.2, 66.5, 65.0, 43.8, 20.0, 18.8; HRMS (EI): Exact mass calcd for C₂₃H₂₃N [M]⁺, 313.1830. Found 313.1838. Relative stereochemistry assigned by NOE experiments (400 MHz NMR).

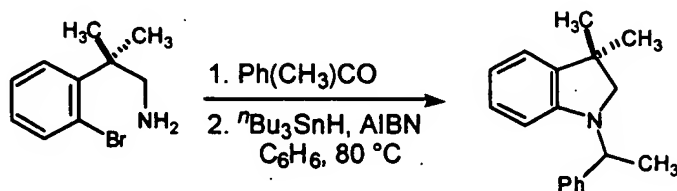
Example 11



[0079] *N*-(1-Phenylbenzyl)-2,3-*cis*-dimethylindoline (5d). Following the general procedure, 2-(*o*-bromophenyl)-1,2-*trans*-dimethylethylamine (18 mg, 79.7 μ mol), benzophenone imine (14.4 mg, 79.7 μ mol) and 4Å MS were stirred in dichloromethane (1 mL) at room temperature for 7.5 days under argon atmosphere to provide the ketimine as an oil; IR (film) 3059, 1624 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.83 (d, J = 7.1 Hz, 2H), 7.62 (dt, J = 7.4, 1.2 Hz, 2H), 7.51 (t, J = 7.7 Hz, 3H), 7.39-7.32 (m, 4H), 7.16 (t, J = 7.3 Hz, 1H), 7.03 (t, J = 7.7 Hz, 1H), 6.80-6.79 (m, 1H), 3.61 (q, J = 6.2 Hz, 1H), 3.54 (q, J = 6.7 Hz, 1H), 1.33 (d, J = 7.0 Hz, 3H), 1.13 (d, J = 6.3 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) ppm 137.2, 132.9, 132.7, 130.3, 129.9, 128.99, 128.5, 128.3, 127.6, 127.3, 61.4, 29.9, 20.6, 16.7; HRMS (EI): Exact mass calcd for C₂₃H₂₂BrN [M+H]⁺, 392.1016. Found 392.1160.

[0080] To a refluxing solution of the unpurified ketimine (5.1 mg, 13 μ mol) in benzene (3 mL) was added n Bu₃SnH (4.6 μ L, 16.9 μ mol), followed by a 6 h addition of AIBN (6.4 mg, 39.0 μ mol) dissolved in benzene (1 mL). After the first 3 h of reaction, another aliquot of n Bu₃SnH (4.6 μ L, 16.9 μ mol), was added to the reaction mixture. After the complete addition of AIBN, the reaction mixture was refluxed for a further 2 h, followed by evaporation of solvent under vacuum. Flash chromatography (5% CH₂Cl₂ in hexanes) yielded the desired product (3.4 mg, 84%) as a gum; IR (film) 3061, 1605 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.45 (d, J = 6.8 Hz, 2H), 7.40 (d, J = 7.2 Hz, 2H), 7.34-7.23 (m, 6H), 7.03 (d, J = 7.1 Hz, 1H), 6.80 (t, J = 7.5 Hz, 1H), 6.63 (t, J = 7.3 Hz, 1H), 5.90 (d, J = 7.8 Hz, 1H), 5.51 (s, 1H), 3.78 (dq, J = 7.5, 6.4 Hz, 1H), 3.38 (dq, J = 7.3, 7.3 Hz, 1H), 1.22 (d, J = 7.1 Hz, 3H), 0.95 (d, J = 6.3 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) ppm 150.0, 142.4, 141.2, 135.5, 128.9, 128.7, 128.6, 127.9, 127.4, 127.2, 126.9, 123.1, 117.5, 109.9, 65.4, 62.4, 39.1, 13.6, 12.6; HRMS (FAB): Exact mass calcd for C₂₃H₂₃N [M]⁺, 313.1830. Found 313.1839.
Relative stereochemistry determined by NOE experiments (400 MHz NMR).

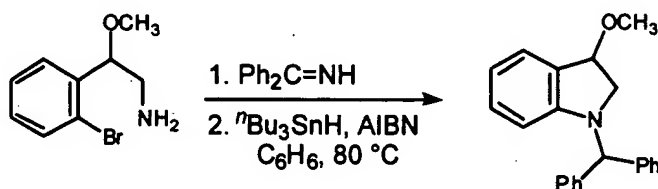
Example 12



[0081] *N*-(1-Methylbenzyl)-3,3-dimethylindoline (5e). According to the general procedure, 2-(*o*-bromophenyl)-2,2-dimethyl ethylamine (733 mg, 3.21 mmol), acetophenone (370 μ L, 3.18 mmol), and 4Å MS were stirred in toluene (25 mL) at room temperature for 12 h to provide the ketimine as a >95:5 mixture of stereoisomers, IR (film) 3058, 1635 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.66 (m, 2H), 7.55 (d, J = 9.1 Hz, 1H), 7.50 (d, J = 9.8 Hz, 1H), 7.30 (t, J = 1.8 Hz, 3H); 7.23 (t, J = 4.3 Hz, 1H), 7.0 (t, J = 9.4 Hz, 1H), 3.98 (s, 2H), 2.20 (s, 3H), 1.64 (s, 6H); ¹³C NMR (100 MHz, CDCl₃) ppm 164.3, 146.3, 141.5, 135.8, 130.0, 129.4, 128.2, 127.7, 127.3, 126.8, 122.8, 60.4, 41.8, 26.8, 26.5, 15.6; HRMS (CI): Exact mass calcd for C₁₈H₂₁BrN, [M+H]⁺, 330.0857. Found 330.0818.

[0082] A three hour addition of $n\text{Bu}_3\text{SnH}$ (72.0 μL , 267 μmol) and AIBN 16 mg, 97 μmol) benzene solution (1.0 mL) to a refluxing solution of the unpurified ketimine (80.2 mg, 243 μmol) in benzene (24 mL) furnished, after flash chromatography (1% Et_2O in hexanes) 48.8 mg (80%) of the desired indoline as a colorless oil. R_f = 0.63 (10% EtOAc /hexanes); IR (film) 3024, 1604 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.41 (d, J = 7.4 Hz, 2H), 7.35 (t, J = 7.2 Hz, 2H), 7.26 (d, J = 7.3 Hz, 1H), 7.01 (t, J = 7.2 Hz, 1H), 7.01 (t, J = 7.2 Hz, 1H), 6.66 (t, J = 8.1 Hz, 1H), 6.35 (d, J = 8.1 Hz, 1H), 4.73 (q, J = 6.9 Hz, 1H), 3.18 (d, J = 8.3 Hz, 1H), 3.08 (d, J = 8.3 Hz, 1H), 1.54 (d, J = 6.9 Hz, 3H), 1.33 (s, 3H), 1.27 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 150.1, 143.4, 139.2, 128.6, 127.5, 127.2, 127.1, 121.8, 117.2, 107.3, 62.6, 54.3, 39.9, 28.3, 27.4, 17.0; HRMS (EI): Exact mass calcd for $\text{C}_{18}\text{H}_{21}\text{N}$ $[\text{M}]^+$, 251.1674. Found 251.1672.

Example 13



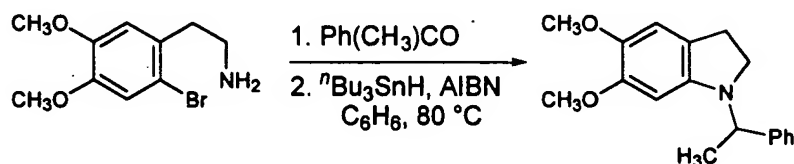
[0083] *N*-(1-Phenylbenzyl)-3-methoxyindoline (5f). 2-(*o*-Bromophenyl)-2-methoxyethylamine (43.4 mg, 189 μmol), benzophenone imine (32 μL , 189 μmol), and 4Å MS were stirred in CH_2Cl_2 at room temperature for 7 h. Removal of the solvent provided the ketimine as a colorless oil. IR (film) 3058, 1626 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.63 (d, J = 8.3 Hz, 2H), 7.52 (dd, J = 8.1, 2.2 Hz, 1H), 7.45–7.28 (m, 8H), 7.13 (dt, J = 9.3, 1.6 Hz, 1H), 7.07 (dd, J = 7.5, 2.4 Hz, 2H), 5.05 (dd, J = 6.7, 4.5 Hz, 1H), 3.71 (dd, J = 14.0, 4.4 Hz, 1H), 3.64 (dd, J = 14.0, 6.9 Hz, 1H), 3.36 (s, 3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 169.8, 140.1 (2C), 137.0, 132.82, 130.2, 129.1, 128.8 (2C), 128.65, 128.6, 128.54, 128.2, 127.7, 123.8, 82.8, 59.2, 57.6; HRMS (EI): Exact mass calcd for $\text{C}_{22}\text{H}_{21}\text{BrNO}$ $[\text{M}+\text{H}]^+$, 394.0808. Found 394.0795.

[0084] A three hour addition of $n\text{Bu}_3\text{SnH}$ (38 μL , 143 μmol) and AIBN (8.5 mg, 52 μmol) in benzene (1 mL) to a refluxing solution of the unpurified ketimine (51.2 mg, 130 μmol) in

benzene (13 mL) afforded, after flash chromatography on basic alumina (10% CH₂Cl₂ in hexanes), 29 mg (70%) of the desired indoline as a colorless oil. R_f = 0.15 (10%

CH₂Cl₂/hexanes); IR (film) 3059, 1608 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.40–7.26 (m, 11H), 7.10 (dt, J = 8.1, 1.2 Hz, 1H), 6.71 (t, J = 7.25 Hz, 1H), 6.34 (d, J = 7.9 Hz, 1H), 5.70 (s, 1H), 4.79 (dd, J = 6.7, 2.6 Hz, 1H), 3.35 (s, 3H), 3.33 (dd, J = 11.3, 2.7 Hz, 1H), 3.27 (dd, J = 11.2, 6.7 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) ppm 140.7, 130.1, 129.2, 128.8, 128.7, 128.4, 128.2, 127.7, 127.5, 126.1, 78.9, 65.7, 56.4, 55.4; HRMS (EI): Exact mass calcd for C₂₂H₂₁NO [M]⁺, 315.1623. Found 315.1623.

Example 14

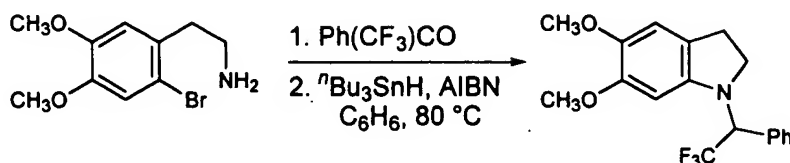


[0085] ***N*-(1-Methylbenzyl)-5,6-dimethoxy indoline (8a).** Following the general procedure, (2-bromo-4,5-dimethoxyphenyl)ethylamine (1.95 g, 7.5 mmol), acetophenone (0.87 mL, 7.5 mmol), and 4Å MS were stirred in toluene (10 mL) at room temperature for 12 h to provide the ketimine as a >95:5 mixture of stereoisomers. IR (film) 3055, 1633 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.77 (m, 2H), 7.38 (m, 3H), 7.03 (s, 1H), 6.83 (s, 1H), 3.87 (s, 3H), 3.79 (s, 3H), 3.77 (dd, J = 7.2, 7.2 Hz, 2H), 3.14 (dd, J = 7.4, 7.4 Hz, 2H), 2.11 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) ppm 166.1, 158.0, 148.3, 148.2, 141.4, 131.9, 129.7, 128.5, 126.7, 115.6, 114.4, 56.3, 56.1, 52.2, 37.3, 15.6; HRMS (CI): Exact mass calcd for C₁₈H₂₁BrNO₂ [MH]⁺, 362.0756. Found 362.0747.

[0086] A three-hour addition of ⁿBu₃SnH (120 μ L), 448 μ mol) and AIBM (26.7 mg, 163 μ mol) solution in benzene (1.1 mL) to a refluxing solution of the unpurified ketimine (148 mg, 408 μ mol) in benzene (41 mL) afforded, after flash chromatography (15% EtOAc in hexanes), 95.6 mg (83%) of the desired indoline as a yellow oil. R_f = 0.27 (20% EtOAc/hexanes); IR (film) 3059, 1614 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.45 (d, J = 7.6 Hz, 2H), 7.36 (t, J = 7.6 Hz, 2H), 7.27 (t, J = 7.4 Hz, 1H), 6.75 (s, 1H), 6.00 (s, 1H), 4.57 (q, J = 6.8 Hz, 1H), 3.80 (s, 3H), 3.70 (s, 3H), 3.34 (dd, J = 16.2, 7.3 Hz, 2H), 2.88 (dd, 7.9, 7.9 Hz, 2H), 1.53 (d, J = 7.0 Hz,

3H); ^{13}C NMR (100 MHz, CDCl_3) ppm 149.0, 146.5, 143.6, 141.5, 128.6, 127.3, 127.1, 121.3, 111.2, 95.4, 57.6, 49.6, 28.4, 17.5; HRMS (EI): Exact mass calcd for $\text{C}_{18}\text{H}_{21}\text{NO}_2$ $[\text{M}]^+$, 283.1572. Found 283.1578.

Example 15



[0087] N-(1-Trifluoromethyl(benzyl))-5,6-dimethoxy indoline (8b). Following the general procedure, (2-bromo-4,5-dimethoxy-phenyl)ethylamine (2.05 g, 7.88 mmol), trifluoroacetophenone (1.1 mL, 7.88 mmol), and 4Å MS were stirred in toluene (30 mL) at room temperature for 12 h to give the ketimine as a >95:5 mixture of stereoisomers. IR (film) 3061, 1669 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.43 (t, $J = 7.2$ Hz, 1H), 7.38 (t, $J = 7.5$ Hz, 2H), 6.95 (s, 1H), 6.91 (d, $J = 7.1$ Hz, 2H), 6.70 (s, 1H), 3.87 (s, 3H), 3.84 (s, 3H), 3.67 (dd, $J = 8.2, 6.8$ Hz, 2H), 3.05 (dd, $J = 6.7, 6.7$ Hz, 2H); ^{13}C NMR (100 MHz, CDCl_3) ppm 159.5, 159.2, 148.5, 148.3, 130.4, 130.2, 130.1, 128.8, 128.5, 127.7, 121.1, 118.4, 115.5, 114.5, 114.2, 56.4, 56.1, 52.9, 36.4; HRMS (EI): Exact mass calcd for $\text{C}_{18}\text{H}_{17}\text{BrF}_3\text{NO}_2$ $[\text{M}]^+$, 415.0395. Found 415.0414.

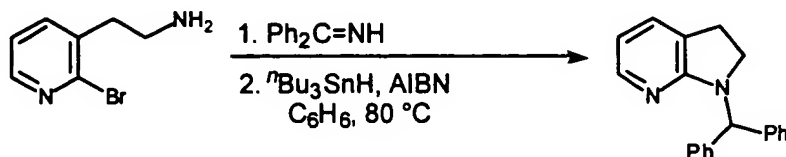
[0088] A three-hour addition of $n\text{Bu}_3\text{SnH}$ (110 μL , 412 μmol) and AIBN (25 mg, 150 μmol) solution in benzene 1.0 mL) to a refluxing solution of the unpurified ketimine (156 mg, 375 μmol) in benzene (37 mL) provided, after flash chromatography (10% EtOAc in hexanes), 113 mg (89%) of the desired indoline as a yellow oil. $R_f = 0.37$ (20% EtOAc/hexanes); IR (film) 2939, 1615 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.39 (m, 5H), 6.74 (s, 1H), 6.27 (s, 1H), 5.11 (q, $J = 8.6$ Hz, 1H), 3.87 (s, 3H), 3.81 (s, 3H), 3.57 (dd, $J = 15.1, 8.8$ Hz, 1H), 3.15 (dd, $J = 18.4, 9.3$ Hz, 1H), 2.89 (m, 2H); ^{13}C NMR (100 MHz, CDCl_3) ppm 149.3, 144.9, 142.4, 132.0, 128.9, 120.2, 111.2, 94.2, 62.8 (q, $J = 29.7$ Hz, 1C), 57.4, 56.6, 49.6, 28.5; HRMS (EI): Exact mass calcd for $\text{C}_{18}\text{H}_{18}\text{F}_3\text{NO}_2$ $[\text{M}]^+$, 337.1290. Found 337.1280.

Example 16

[0089] *N*-(1-Phenylbenzyl)-5,6-dimethoxy indoline (8c). Following the general procedure, (2-bromo-4,5-dimethoxyphenyl)ethyl amine (343 mg, 1.32 mmol) and benzophenone imine (239 mg, 1.32 mmol) were stirred at room temperature in CH_2Cl_2 (3 mL) for 12 h to give the ketimine. IR (film) 3056, 1623 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.62 (d, $J = 7.0$ Hz, 2H), 7.40 (dd, $J = 6.2, 3.4$ Hz, 3H), 7.36 (t, $J = 7.6$ Hz, 3H), 6.95 (s, 1H), 6.94 (d, $J = 2.3$ Hz, 2H), 6.73 (s, 1H), 3.84 (s, 3H), 3.76 (s, 3H), 3.67 (dd, $J = 7.0, 7.0$ Hz, 2H), 3.07 (dd, $J = 7.2, 7.2$ Hz, 2H); ^{13}C NMR (100 MHz, CDCl_3) ppm 168.9, 148.2, 139.9, 136.8, 131.8, 128.5, 128.4, 128.3, 127.9, 115.5, 114.5, 114.2, 56.3, 56.1, 53.8, 37.4; HRMS (CI): Exact mass calcd for $\text{C}_{23}\text{H}_{23}\text{BrNO}_2$ $[\text{M}+\text{H}]^+$, 424.0912. Found 424.0910.

[0090] A three-hour addition of $n\text{Bu}_3\text{SnH}$ (75 μL , 278 μmol) and AIBN (17 mg, 101 μmol) solution in benzene (1.0 mL) to refluxing solution of the unpurified ketimine (107 mg, 252 μmol) in benzene (25 mL) provided, after flash chromatography (5% EtOAc in hexanes), 46 mg (64%) of the desired indoline as a white solid. mp 105–106 $^\circ\text{C}$; $R_f = 0.20$ (10% EtOAc/hexanes); IR (film) 2931, 1597 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.45 (d, $J = 7.1$ Hz, 4H), 7.35 (t, $J = 7.1$ Hz, 4H), 7.27 (t, $J = 4.7$ Hz, 2H), 6.75 (s, 1H), 5.79 (s, 1H), 5.33 (s, 1H), 3.80 (s, 3H), 3.51 (s, 3H), 3.16 (dd, $J = 8.3, 8.3$ Hz, 2H), 2.87 (dd, $J = 8.1, 8.1$ Hz, 2H); ^{13}C NMR (100 MHz, CDCl_3) ppm 148.4, 146.9, 142.1, 141.8, 128.7, 128.5, 127.4, 121.7, 110.4, 96.5, 69.3, 57.3, 55.9, 53.6, 28.5; HRMS (CI): Exact mass calcd for $\text{C}_{23}\text{H}_{23}\text{NO}_2$ $[\text{M}]^+$, 345.1729. Found 345.1713.

Example 17



[0091] *N*-(1-Phenylbenzyl)-6-azaindoline (8d). 2-Bromo-3-(2-aminoethyl) pyridine (104 mg, 517 μmol) and benzophenone imine (94 mg, 517 μmol) were stirred in CH_2Cl_2 at room temperature for 8 h to provide the ketimine. IR (film) 3056, 1622 cm^{-1} ; ^1H NMR (400 MHz,

CDCl₃) δ 8.20 (dd, J = 4.7, 0.9 Hz, 1H), 7.56 (t, J = 7.8 Hz, 3H), 7.43–7.40 (m, 3H), 7.37 (d, J = 6.6 Hz, 1H), 7.32 (t, J = 7.9 Hz, 2H), 7.16 (dd, J = 7.4, 5.2 Hz, 1H); 6.96 (t, J = 3.4 Hz, 2H), 3.69 (t, J = 6.9 Hz, 2H), 3.10 (t, J = 6.9 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) ppm 169.2, 158.1, 147.7, 144.5, 139.5, 137.0, 136.5, 130.0, 128.5, 128.4, 128.3, 127.5, 122.6, 52.4, 36.8; HRMS (EI): Exact mass calcd for C₂₀H₁₇BrN₂ [M]⁺, 364.0575. Found 364.0587.

[0092] A two-hour addition of ⁿBu₃SnH (58 μ L, 214 μ mol) and AIBN (39 mg, 235 μ mol) solution in benzene (2 mL) to a refluxing solution of the unpurified ketimine (36 mg, 98 μ mol) in benzene (10 mL) delivered, after flash chromatography (5% EtOAc/Hexanes), 14 mg (50%) of the desired indoline as an orange crystalline solid. mp 101 °C; R_f = 0.1 (5% EtOAc/Hexanes); IR (film) 3058, 1611 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 7.87 (d, J = 5.2 Hz, 1H), 7.34–7.26 (m, 10H), 7.17 (d, J = 7.0 Hz, 1H), 6.81 (s, 1H), 6.42 (t, J = 6.5 Hz, 1H), 3.34 (t, J = 8.5 Hz, 2H), 2.97 (t, J = 8.5 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) ppm 162.8, 146.1, 140.2, 131.2, 122.8, 129.1, 128.5, 127.4, 112.4, 60.0, 45.6, 25.9; HRMS (EI) Calcd for C₂₀H₁₈N₂ [M]⁺, 286.1470. Found 286.1468.

Example 33

General Example for Vinyl Amination

[0093] 4-Pentyn-1-amine (0.59g) was dissolved in 8mL of dry benzene, and 4A molecular sieves (~4 g) were added to the solution. The mixture was stirred at room temperature and 0.78 mL of acetophenone was added dropwise. The mixture was allowed to stir for at least 6 hours. The reaction mixture was filtered through a plug of celite, and the molecular sieves were washed with ether. The solvent was washed *in vacuo* to afford the 4-pentyn-1-imine in 94% yield.

[0094] The imine (0.1709 g) was dissolved in 46 mL benzene and heated to reflux at 85°. AIBN (0.0224 g, 0.2 eq.) and tri-*n*-butyltin hydride (0.25 mL, eq.) were dissolved in a minimal amount of benzene (~1 mL) and drawn into syringe. This solution was added dropwise over 3 hours via syringe pump to the refluxing imine solution. After complete addition of AIBN and tri-*n*-butyltin hydride, the mixture was allowed to reflux for an additional 3 hours. Once the reaction had gone to completion, the reaction mixture was cooled to room temperature and the solvent was removed *in vacuo*.